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UNITED STATES ARMY ENVIRONMENTAL HYGIENE AGENCÝ

aberdeen proving Ground, MD 21019

TOPICAL HAZARD EVALUATION PROGRAM

OF

CANDIDATE INSECT REPELLENTS

AI3-38221a and AI3-38351a

US DEPARTMENT OF AGRICULTURE PROPRIETARY CHEMICALS

STUDY NOS. 75-51-0333-84 and 75-51-0370-84

JUNE 1981 - DECEMBER 1983



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Eye Irritation USDA Proprietary Chem	_			
Guinea Pig Sensitization	ilcais			
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DEPARTMENT OF THE ARMY U. S. ARMY ENVIRONMENTAL HYGIENE AGENCY ABERDEEN PROVING GROUND, MARYLAND 21010

REPLY TO ATTENTION OF

HSHB-OT/WP

3 APR 1001

SUBJECT: Topical Hazard Evaluation Program of Candidate Insect Repellents

AI3-38221a and AI3-38351a, US Department of Agriculture Proprietary Chemicals, Study Nos. 75-51-0333-84 and

75-51-0370-84, June 1981 - December 1983

Executive Secretary Armed Forces Pest Management Board Forest Glen Section, WRAMC Washington, DC 20307

EXECUTIVE SUMMARY

The purpose, essential findings, and major recommendations of the inclosed report follow:

- a. <u>Purpose</u>. The purpose of this program is to provide guidance for further entomological testing of the candidate insect repellents AI3-38221a and AI3-38351a by means of laboratory animal studies using New Zealand White rabbits, Sprague-Dawley rats, and albino Hartley guinea pigs.
- b. <u>Essential Findings</u>. Chemical AI3-38221a produced mild primary irritation of the intact skin and of the skin surrounding an abrasion. Chemical AI3-38351a produced no primary irritation of the intact skin and no greater than mild primary irritation of the skin surrounding an abrasion. Both chemicals produced mild injury to the cornea and, in addition, some injury to the conjunctiva. Chemicals AI3-38221a and AI3-38351a did not produce sensitization or photoirritation reactions and were relatively nontoxic by ingestion.
- c. <u>Major Recommendations</u>. Recommend that chemicals AI3-3822la and AI3-3835la be approved for further testing as candidate insect repellents.

FOR THE COMMANDER:

l Incl as (5 cy) Director, Occupational and
Environmental Health

CF:
HQDA (DASG-PSP) wo incl
Cdr, HSC (HSCL-P)
Comdt, AHS (HSHA-IPM)
Dir, Advisory Cen on TOX, NRC (2 cy)
USDA, ARS (Dr. Terrence McGovern)
USDA, ARS-Southern Region (3 cy)
Cdr, USAMRDC [SGRD-DPM/LTC(P) Reinert]



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DEPARTMENT OF THE ARMY U. S. ARMY ENVIRONMENTAL HYGIENE AGENCY ABERDEEN PROVING GROUND, MARYLAND 21010

TOPICAL HAZARD EVALUATION PROGRAM OF

CANDIDATE INSECT REPELLENTS
AI3-38221a and AI3-38351a
US DEPARTMENT OF AGRICULTURE PROPRIETARY CHEMICALS
STUDY NOS. 75-51-0333-84 and 75-51-0370-84
JUNE 1981 - DECEMBER 1983

1. AUTHORITY.

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- a. Letter, US Department of Agriculture Agricultural Research, Southern Region, Insects Affecting Man and Animals Research Laboratory, Gainesville, Florida, 18 June 1981.
- b. Letter, US Department of Agriculture Agricultural Research, Southern Region, Insects Affecting Man and Animals Research Laboratory, Gainesville, Florida, 29 April 1982.
- c. Memorandum of Understanding between the US Army Environmental Hygiene Agency; the US Army Health Services Command; the Department of the Army, Office of The Surgeon General; the Armed Forces Pest Control Board; and the Department of Agriculture, Agriculture Research, Science and Education Administrations, titled Coordination of Biological and Toxicological Testing of Pesticides, effective 23 January 1979.
- 2. REFERENCE. Toxicology Division Topical Hazard Evaluation Program Procedural Guide, US Army Environmental Hygiene Agency (USAEHA), January 1982.
- 3. PURPOSE. The purpose of this program is to provide guidance for further entomological testing of the candidate insect repellents AI3-38221a and AI3-38351a, US Department of Agriculture (USDA) Proprietary Chemicals.
- 4. SUMMARY OF FINDINGS. Hazard evaluations of the candidate insect repellents AI3-38221a and AI3-38351a, USDA Proprietary Chemicals, were conducted by this Agency using New Zealand White rabbits, Sprague-Dawley rats, and albino Hartley guinea pigs. A tabular presentation of animal toxicity data developed by this Agency follows:

† The studies reported herein were performed in animal facilities fully accredited by the American Association for the Accreditation of Laboratory Animal Care.

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^{*} In conducting the studies described in this report, the investigators adhered to the "Guide for the Care and Use of Laboratory Animals," US Department of Health, Education, and Helfare; Public Health Service; National Institutes of Health Publication No. 80-23, revised 1978, reprinted April 1980.

TABLE. PRESENTATION OF DATA

AND THE STREET SECRET SECRET SECRET SECRET SECRET SECRET SECRET SECRET.

,TEST	RESUL	rs	INTERPRETATION	
SKIN IRRITATION STUDIES				
Rabbits				
to intact and abraded skin pro of New Zealand White rabbits. irr int 0.5 mL technical grade ski	Chemical Alproduced mi irritation intact skir surrouabrasion.	ld prfmary of the and of the	USAEHA Category II (ref Appendix A)	
	Chemical AI produced no irritation intact skin more than mirritation skin surrouabrasion.	primary of the and no ild of the	USAEHA Category I (ref Appendix A)	
EYE IRRITATION STUDIES				
Rabbits				
Single 24-hour application of 0.1 mL technical grade chemical to one eye of each of nine New Zealand White rabbits. Three of the nine rabbits had the eye flushed with warm water for 1 minute, 25 seconds after application.	mild injury	5la produced to the in addition, to the	USAEHA Category C (ref Appendix A)	
APPROXIMATE LETHAL DOSE				
Oral Rats (male) – no diluent	AI3-38221a	2,870mg/kg	These chemicals are	
Rats (female) - no diluent	AI3-38351a	3,330mg/kg	relatively nontoxic by ingestion.	

TEST

RESULTS

INTERPRETATION

PHOTOCHEMICAL SKIN IRRITATION STUDIES

Rabbits

A single 0.05 mL application of a 25% (w/v) solution of each chemical and of a 10% (w/v) Oil of Bergamot solution (positive control) in 95% ethyl alcohol was applied to the intact skin of six rabbits. Five minutes after application, the rabbits were exposed to ultraviolet (UV) light (365 nm) for 30 minutes at a distance of 10-15 cm.

These chemicals did not produce photochemical irritation under test conditions. These chemicals did not produce photo-chemical irritation under test conditions and are not expected to produce photo-irritation in humans.

Control

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Following UV exposure of the rabbits, 0.05 mL of test chemicals, positive control and diluent were applied to additional skin areas to serve as unirradiated control sites. Application areas were checked for skin irritation at 24, 48, and 72 hours.

SENSITIZATION STUDIES

Guinea Pigs (Female)

Intradermal (ID) injections of 0.1 mL of a minimally irritating concentration of each test chemical or of dinitrochlorobenzene (DNCB)* in a mixture containing 1 volume of propylene glycol and 29 volumes of saline.

Positive control application and irradiation caused greater irritant effects than on unirradiated skin areas.

^{*} A known skin sensitizer.

TEST	ST RESULTS	
duines nigs for	Challenge doses of	These chemicals a

Ten test guinea pigs for each chemical were given 10 sensitizing doses over a 3-week period. After a 2-week rest, they were challenged with ID injections of each test chemical.

Challenge doses of chemicals AI3-39221a and AI3-38351a did not produce a sensitization reaction.

These chemicals are not expected to produce a sensitization reaction in humans.

Control

Ten positive control guinea pigs were sensitized over 3 weeks with DNCB.
After a 2-week rest, they were challenged with ID injections of DNCB.

Challenge dose of DNCB in positive control guinea pigs produced a marked sensitization reaction in 10 out of 10 guinea pigs.

DNCB produced a marked sensitization reaction, indicating that these guinea pigs respond to sensitizing agents.

- 5. CONCLUSION. Chemical AI3-38221a produced mild primary irritation of the intact skin and of the skin surrounding an abrasion. Chemical AI3-38351a produced no primary irritation of the intact skin and no greater than mild primary irritation of the skin surrounding on abrasion. Both chemicals produced mild injury to the cornea and, in addition, some injury to the conjunctiva. Chemicals AI3-38221a and AI3-38351a did not produce sensitization or photoirritation reactions and were relatively nontoxic by ingestion. These studies were monitored by Analytical Quality Assurance Office (see Appendix B).
- 6. RECOMMENDATION. Recommend that chemicals AI3-38221a and AI3-38351a be approved for further testing as candidate insect repellents.

JOHN V. WADE, DVM

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Laboratory Animal Veterinary Officer

Toxicology Division

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APPROVED:

MAURICE H. HEEKS

Chief, Toxicology Division

APPENDIX A

TOPICAL HAZARD EVALUATION PROGRAM DEFINITIONS OF CATEGORIES OF COMPOUNDS BEING CONSIDERED FOR ACUTE SKIN APPLICATION

- <u>CATEGORY I</u> Compounds producing no primary irritation of the intact skin or no greater than mild primary irritation of the skin surrounding an abrasion. (INTERPRETATION: No restriction for acute application to the human skin.)
- CATEGORY II Compounds producing mild primary irritation of the intact skin and the skin surrounding an abrasion. (INTERPRETATION: Should be used only on human skin found by examination to have no abrasions or may be used as a clothing impregnant.)
- CATEGORY III Compounds producing moderate primary irritation of the intact skin and the skin surrounding an abrasion. (INTERPRETATION: Should not be used directly on the skin without a prophetic patch test having been conducted on humans to determine irritation potential to human skin. May be used without patch testing, with extreme caution, as clothing impregnants. Compound should be resubmitted in the form and at the intended use concentration so that its irritation potential can be reexamined using other test techniques on animals.)
- CATEGORY IV Compounds producing moderate to severe primary irritation of the intact skin and of the skin surrounding an abrasion and, in addition, producing necrosis, vesiculation, and/or eschars. (INTERPRETATION: Should be resubmitted for testing in the form and at the intended use concentration. Upon resubmission, its irritation potential will be reexamined using other test techniques on animals, prior to possible prophetic patch testing in humans, at concentrations which have been shown not to produce primary irritation in animals.)
- CATEGORY V Compounds impossible to classify because of staining of the skin or other masking effects owing to physical properties of the compound. (INTERPRETATION: Not suitable for use on humans.)

EYE CATEGORIES:

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- A. <u>Compounds noninjurious to the eye</u>. INTERPRETATION: Irritation of human eyes is not expected if the compound should accidentally get into the eyes, provided it is washed out as soon as possible.
- B. <u>Compounds producing mild injury to the cornea</u>. INTERPRETATION: Should be used with caution around the eyes.
- C. Compounds producing mild injury to the cornea, and in addition some injury to the conjunctiva. INTERPRETATION: Should be used with caution around the eyes and mucosa.
- D. <u>Compounds producing moderate injury to the cornea</u>. INTERPRETATION: Should be used with extreme caution around the eyes.
- E. Compounds producing moderate injury to the cornea, and in addition producing some injury to the conjunctiva. INTERPRETATION: Should be used with extreme caution around the eyes and mucosa.
- F. Compounds producing severe injury to the cornea and to the conjunctiva. INTERPRETATION: Should be used with extreme caution. It is recommended that use be restricted to areas other than the face.

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APPENDIX B

ANALYTICAL QUALITY ASSURANCE

The Analytical Quality Assurance Office certifies the following:

a. These studies were conducted in accordance with:

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- (1) Standing Operating Procedures developed by the Toxicology Division, USAEHA.
- (2) Title 21, Code of Federal Regulations (CFR), 1983 rev, Part 58, Good Laboratory Practice for Nonclinical Laboratory Studies.
- (3) Final Rule, Pesticide Programs; Good Laboratory Practice Standards; 48 Federal Register (FR) 53963-539691, 29 November 1983.
- b. Facilities were inspected during its operational phase to ensure compliance with paragraph a above.
- c. The information presented in this report accurately reflects the raw data generated during the course of conducting these studies.

PAUL V. SNEERINGER, Ph.D. Chief, Analytical Quality Assurance Office